2003 2004

PCT/US2004/040872

PATENT COOPERATION TREATY

NTERNATIONAL SEARCE	TING AUTHORITY		•			
To: DANIEL HART KNOBBB, MARTENS, OLSON & BEAR, LLP			PCT			
2040 MAIN STREET			WRI	TTEN OPINION OF THE		
14TH FLOOR IRVINE, CA 92614		ļ	INTERNATIO	NAL SEARCHING AUTHOR	UTY	
				(PCT Rule 43bis.1)	·	
		،بدل	Date of mailing (day/month/year)	13 NOV 2006		
Applicant's or agent's file r	eference		FOR FURTHER ACTION See paragraph 2 below			
LBNL.001VPC International application No	Tritarnat	onal filing date	(day/month/year)	Priority date (day/month/year)		
		mber 2004 (06.	•	04 December 2003 (04.12.2003)		
PCT/US04/40872 International Patent Classifi	igntion (IPC) or both na	tional classificat	ion and IPC			
	06.01) C12Q 1/68(
Applicant .						
REGENTS OF THE UNIV	ersity of Califor	NIA				
1. This opinion contains	indications relating to th	ne following item	19!			
Box No. 1	Basis of the opinion					
Box No, Il	Priority	,				
Box No. III	Box No. III Non-establishment of opinion with regard to nevelty, inventive step and industrial applicability					
Box No. IV	Lack of unity of inve	ention			1	
Box No. V	Reasoned statement applicability; citation	Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
Box No. VI	Certain documents of					
Box No. VII	Cortain defects in the	o international a	pplication	•	}	
Box No. VIII	Certain observations	on the internati	onal application			
Authority other than that written opinions of	national preliminary ex nary Examining Author this one to be the IPBA of this international Sca	and the choser rehing Authority	PEA has notified the will not be so consider		6.1 <i>bls(b</i>)	
of Form PCT/ISA/226	together, where approp or before the expiration	lered to be a wr riste, with amen on of 22 months	itten opinion of the II denents, before the ex from the priority date,	PBA, the applicant is invited to sub- piration of 3 manths from the date of whichever expires later.	nit to the finalling	
For further options, se	e Form PCT/ISA/220.					
3, For further details, se	s notes to Form PCT/IS	A/220.				
Name and mailing address	s of the ISA/US	Data of comp	letion of this opinion	Authorized officer Both	Halle	
Mail Stop PCT, A	un: ISA/US	1	006 (01.10,2006)	M. Franco Salvoza	Harris	
Commissioner for P.O. Box 1450 Alexandria, Visgi		0.00000	V. M (Telephone No. (571) 272-1640	9	
Facsimile No. (571) 273-3 Form PCT/ISA/237 (cover	3201	<u> </u>				

International application No.
PCT/US04/40872

Box No	o. I Basis of this opinion					
. With	regard to the language, this opinion has been established on the basis of:	,				
\boxtimes	the international application in the language in which it was filed					
	a translation of the international application into, which is the language of a translation furnished international search (Rules 12.3(a) and 23.1(b)).	. 1				
2. With inven	regard to any nucleotide and/or amino acid sequence disclosed in the international application and nece tion, this opinion has been established on the basis of:	ssary to the claimed				
a.	type of material					
	a sequence listing					
	table(s) related to the sequence listing					
ъ.	format of material					
	on paper					
	in electronic form					
c.	time of filing/furnishing					
	contained in the international application as filed.					
	filed together with the international application in electronic form.					
	furnished subsequently to this Authority for the purposes of search.	1				
	Turnished subsequently to this realistic, to the party of					
3	In addition, in the case that more than one version or copy of a sequence listing and/or table(s) relating or furnished, the required statements that the information in the subsequent or additional copies is application as filed or does not go beyond the application as filed, as appropriate, were furnished.	thereto has been filed dentical to that in the				
4. Add	litional comments:					
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Form PCT/ISA/237(Box No. I) (April 2005)

International application No. PCT/US04/40872

Box No. V Reasoned statement under Rule 43 bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement					
Statement					
Novelty (N)		Please See Continuation Sheet	YES		
	Claims	Please See Continuation Sheet	ио		
	Claima	Please See Continuation Sheet			
Inventive step (IS)		Please See Continuation Sheet	YES NO		
	V.a	, INNO 435 SELLING			
Industrial applicability (IA)	Claims	s Please See Continuation Sheet			
	Claims	Please See Continuation Sheet	ио		
Citations and explanations:					
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Box No.	VII	Certain	defects in	the international	application
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The following defects in the form or contents of the international application have been noted:

Claim 30 is objected to under PCT Rule 66.2(a)(iii) as containing the following defect(s) in the form or contents thereof: It contains a misspelling of the term "florescence."

Form PCT/ISA/237 (Box No. VII) (April 2005)

International application No. PCT/US04/40872

Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

V.1. Reasoned Statements:

The opinion as to Novelty was positive (Yes) with respect to claims 2-6, 9, 12, 15-17, 19-24, 29, 31

The opinion as to Novelty was negative (No) with respect to claims 1, 7, 8, 10, 11, 13, 14, 18, 25, 26, 27, 28, 30, 32

The opinion as to Inventive Step was positive (Yes) with respect to claims NONE
The opinion as to Inventive Step was negative(NO) with respect to claims 1, 7, 8, 10, 11, 13, 14, 18, 25, 26, 27, 28, 30, 32

The opinion as to Industrial Applicability was positive (YES) with respect to claims 1-32

The opinion as to Industrial Applicability was negative(NO) with respect to claims NONE

V. 2. Citations and Explanations:

Claims 1, 7, 8, 10, 11, 18, 25, 28, 30, 32 novelty under PCT Article 33(2) as being anticipated by TANG et al. (2001).

Claim I recites a method for detecting an analyte in a sample, comprising: providing a suspension of colloidal particles, wherein said particles are associated with a ligand that binds to said analyte, and wherein said colloidal particles are near a dynamical phase transition state; contacting said suspension with said sample; and determining whether said colloidal particles transition from a first phase to a second phase, wherein such transition is indicative of said analyte being present in said sample.

Claims 7, 8, 10, 11 further recite the method of claim 1 wherein said ligand is non-covalently linked to said colloidal particles; wherein said ligand is interspersed within a lipid layer on said colloidal particles; wherein said analyte is selected from the group consisting of a protein, a nucleic acid, an antibody, an antigen, a receptor, a virus, and a bacteria; wherein determining whether said colloidal particles transition from a first phase to a second phase comprises measuring the distances between centers of said colloidal particles in said suspension.

Claims 13, 14 recite the method of claim 1, wherein said first phase is a condensed phase and said second phase is a dispersed phase; wherein said first phase is a dispersed phase and said second phase is a condensed phase.

Claim 18 recites an assay system for detecting the binding, comprising: a suspension of colloidal particles, wherein said particles are near a dynamical phase transition state; a ligand associated with said particles and specific for said analyte; and a device configured to determine if said colloidal particles transition from a first phase to a second phase when contacted by said analyte, wherein such transition is indicative of said analyte being present in said sample.

Claim 25 further recites the system of claim 18 wherein said ligand is non-covalently linked to said colloidal particles.

Claims 13, 14 recite the system of claim 18, wherein said first phase is a condensed phase and said second phase is a dispersed phase; wherein said first phase is a dispersed phase and said second phase is a condensed phase.

Claim 28 recites an assay system as recited above further comprising a means for detecting is said colloidal particles transition from a first phase to a second phase when contacted by said analyte, wherein such transition is indicative of said analyte being bound to said ligand

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Supplemental Box In case the space in any of the preceding boxes is not sufficient.

Claims 30, 32 further recite the system of claim 28 wherein said means comprises a florescence detector; wherein said ligand is non-covalently linked to said colloidal particles.

TANG et al. teaches providing a suspension of DNA winked colloidal nanoparticles above the phase transition temperature of the polyNIPAAm part (p. 165). A complementary ODN was added to the dispersion, wherein the particles dispersed in the absence of the complementary ODN, and aggregated in the presence of the complementary DNA. The analyte is a nucleic acid as recited in claim 10; the distances between the colloidal particles was measured as they were measured in a dispersed phase as opposed to an aggregated one. Further, a decrease in transmittance was measured in the conjugate solution containing the complementary ODN. Thus, TANG et al. teaches a method and device determining that the DNA-linked colloidal nanoparticles aggregate depending on the DNA hybridization (p. 166).

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